



**UNITED STATES DEPARTMENT OF COMMERCE**  
**Patent and Trademark Office**

Address: COMMISSIONER OF PATENTS AND TRADEMARKS  
Washington, D.C. 20231

*Pol*

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.
-----------------	-------------	----------------------	---------------------

08/913,918 12/08/97 PROCKOP

D TJU-1857

EXAMINER
----------

000570 HM12/1012  
AKIN, GUMP, STRAUSS, HAUER & FELD, L.L.P.  
ONE COMMERCE SQUARE  
2005 MARKET STREET, SUITE 2200  
PHILADELPHIA PA 19103

NGUYEN, D	
ART UNIT	PAPER NUMBER

1633

*34*

DATE MAILED:

10/12/01

**Please find below and/or attached an Office communication concerning this application or proceeding.**

**Commissioner of Patents and Trademarks**

# Office Action Summary

Application No.

08/913,918

Applicant(s)

PROCKOP ET AL.

Examiner

Dave Nguyen

Art Unit

1633

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

## Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

## Status

- 1) ☐ Responsive to communication(s) filed on \_\_\_\_.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

## Disposition of Claims

- 4) ☒ Claim(s) 69-112 is/are pending in the application.
- 4a) Of the above claim(s) \_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 101-112 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_ is/are objected to.
- 8) ☒ Claim(s) 69-100 are subject to restriction and/or election requirement.

## Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
- Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- 11) ☐ The proposed drawing correction filed on \_\_\_\_ is: a) ☐ approved b) ☐ disapproved by the Examiner.
- If approved, corrected drawings are required in reply to this Office action.
- 12) ☐ The oath or declaration is objected to by the Examiner.

## Priority under 35 U.S.C. §§ 119 and 120

- 13) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some \* c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- \* See the attached detailed Office action for a list of the certified copies not received.
- 14) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).
- a) ☐ The translation of the foreign language provisional application has been received.
- 15) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

## Attachment(s)

- 1) ☒ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☒ Information Disclosure Statement(s) (PTO-1449) Paper No(s) 24
- 4) ☒ Interview Summary (PTO-413) Paper No(s) 24
- 5) ☐ Notice of Informal Patent Application (PTO-152)
- 6) ☐ Other:

Art Unit: 1633

This application has been assigned to a new examiner since the previous examiner has left the Office. The request filed 12/20/00 for a Continued Prosecution Application (CPA) under 37 CFR 1.53(d) based on parent Application No 08/913,918 is acceptable and a CPA has been established. An action on the CPA follows. Claims 1-39 have been canceled; Claims 40-43 have been amended; and claim 44 has been added by the preliminary amendment filed on July 9, 1999.

Prosecution of the previously elected invention is being continued in this CPA since Applicants did not indicate a desire to elect a different invention for examination, MPEP 819.

As the result, given Applicant's election of the species of obesity factor in Paper No. 13, filed 9/15/99, claims that do not embrace the elected species, e.g., claims 76, 78-96, are withdrawn from further consideration by the Examiner, 37 C.F.R. § 1.142(b), as being drawn to a non-elected species.

In addition, Applicant's election without traverse of species of osteoporosis (as recited in claim 75) in the response filed July 16, 2001 is acknowledged.

Thus, elected claims 69-75 and 97-112 readable on the species of osteoporosis and obesity factor, to which following grounds of rejection are applicable, are pending for examination.

35 U.S.C. 101 reads as follows:

Whoever invents or discovers any new and useful process, machine, manufacture, or composition of matter or any new and useful improvement thereof, may obtain a patent therefore, subject to the conditions and requirements of this title.

Claims 72-75 are rejected under 35 USC 101 because the claims are readable on an animal and/or human comprising the implanted container of claim 69. A claim including within its scope a human being is not considered patentable subject matter as the limited but exclusive property right in a human being is barred by the United States Constitution. See 1077 OG 24.

In addition, the as-filed specification does not provide any credible, substantial and specific utility of the claimed animal or human. A disclosed utility of the container of claim 69 for expression of recombinant proteins and/or for medical uses in an animal is not the same as the lack credible, substantial and specific utility of the claimed animal or human.

Art Unit: 1633

***Claim Rejections - 35 USC § 112***

The following is a quotation of the first paragraph of 35 U.S.C. § 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 73-75, 109-112 are rejected under 35 U.S.C. 112, first paragraph, because the specification is enabling only for claims limited to:

A method of providing a protein to a mammal, the method comprising implanting within the animal a container containing an isolated marrow stromal cell which comprises an expressible gene construct encoding the protein, wherein the container physically isolates the stromal cell from immune cells of the animal, and wherein the container has pores which permit diffusion of the protein between the interior and exterior of the container.

The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to use the invention commensurate in scope with these claims readable on embodiment within the context of *ex vivo* gene therapy of using any genetically modified stromal cells expressing a therapeutic DNA to treat any animal afflicted with a disease, disorder, or condition characterized by a defect in said DNA in the animal.

Factors to be considered in determining whether a disclosure would require undue experimentation have been summarized in In re Wands, 858 F.2d 731, 8USPQ2d 1400 (Fed. Cir. 1988). They include (1) the quantity of experimentation necessary, (2) the amount of direction or guidance presented, (3) the presence or absence of working examples, (4) the nature of the invention, (5) the state of the prior art, (6) the relative skill of those in the art, (7) the predictability or unpredictability of the art, and (8) the breadth of the claims.

Art Unit: 1633

With respect to claims 72-75 readable on animals or humans containing the container of claim 69, the application does not provide any reasonable enablement for a skilled artisan to make and use the claimed animal, particularly in view of the reasons set forth in the 101 rejection.

With respect to claims 72-75 that may be readable on the intended use of the container of claim 69 to treat any animal having any condition, disease, or disorder associated with a defect of a protein, the application and claims 109-111 contemplates that by using any container that contain isolated stromal cells expressing a therapeutic protein encoded DNA in an *ex vivo* gene therapy method in any animal including reptiles, birds, amphibians, mammals and humans having a defect in the protein expressed by the encoded DNA, any condition, disease, or disorder associated with the defect in the animal will be treated therapeutically. Claim 75 lists a number of diseases or disorders including osteoporosis that can be treated by any of the disclosed container having isolated and genetically modified stromal cells incorporated therein.

The application indicates that implantation of mesenchymal cells from normal mice into irradiated transgenic mice that expresses the mutated COL1 A1 gene led to production of progeny cells that express the normal pro $\alpha$ (1) chains in the irradiated mice. The application further contemplates that long-term expression of any therapeutic protein encoded DNA including the obesity gene (Ob) can be achieved for therapeutic application of the gene in the treatments of obesity or decrease of appetite in any animal by using any autologous or non-autologous stromal cell (not necessarily limited to mesenchymal cells (MSC)) having any Ob expressing DNA vector incorporated therein.

While the application including the state of the prior art provides sufficient guidance and reasonable enablement for an improved methods of using claimed container to express a recombinant protein for non-therapeutic applications that are well known to a skilled artisan at the time the invention was made, the application on the basis of applicant's disclosure does not provide any reasonable enablement including sufficient guidance and/or factual evidence so as to reasonably extrapolate from a simple production of endogenous COL1 A1 proteins by allogeneic MSC in transgenic mice expressing the mutated COL1 A1 protein to any therapeutically relevant effect in any animal having a real-world medical disease associated

Art Unit: 1633

with a protein defect, particularly given that *ex vivo* gene therapy of using bone marrow stromal, stromal stem cells or mesenchymal stem cells is an emerging technology that remains reasonably unpredictable at the time the invention was made (see Marshall (Science, Vol. 269, pp. 1050, 1995), Verma *et al.* (Nature, Vol. 389, pp. 239-242), Anderson (Nature, Vol. 292, 25-30, 1998), Moritz *et al.* (J. Clin. Invest. 1994, 93:1451-1457), Riddell *et al.* (Nature Medicine, Vol. 2, 2:216-223, 1996), Onodera *et al.*, Acta Haematologica, 101, 2, pp. 89-96, 1999, Kohn, Current Opinion in Pediatrics, 7, 56-63, 1995).

The specification does not provide reasonable enablement for claims encompassing *ex vivo* gene therapy methods as claimed, wherein any administration route is employed, wherein any genetically modified stromal stem cell is employed, and wherein any disease as disclosed in the claims is contemplated.

More specifically as to the state of the art of *ex vivo* gene therapy of using bone marrow stromal cells including stromal stem cells, the state of the art exemplified by Marshall (Science, Vol. 269, pp. 1050, 1995) indicates that in 1995, "so far, there has been no unambiguous evidence that genetic treatment has produced therapeutic benefits" and that "while there are several reports of convincing gene transfer and expression, there is still little or no evidence of therapeutic benefit in patient or even in animal models" (page 1050). More specifically as the unpredictability of using retroviral vectors which has been preferably used in experimental protocols *in vitro* and/or *in vivo*, due to its ability to better express a recombinant protein in stem cells as compared to other viral or non-viral vectors, Verma *et al.* (Nature, Vol. 389, pp. 239-242) teach that "another formidable challenge to the *ex vivo* approach is the efficiency of transplantation of the infected cells" and that "successful animal models will prove inadequate when the same protocols are extended to humans" (page 240, column 3, last paragraph). The specification, for example, contemplates that by employing any retroviral transduced MSC as carrier of protein drugs in any *ex vivo* gene therapy method, the stem cells would function as a continuous supply of protein drugs to any target site *in vivo*, such that any disease or disorder can be treated therapeutically in any animal. A skilled artisan, attempting to make and use the claimed constructs, would first look to the specification for guidance as to which

Art Unit: 1633

therapeutic protein drug encoded DNA to use in an *ex vivo* gene therapy wherein retroviral transduced bone marrow stromal or stromal stem cells are employed for treating a disorder. However, the state of the prior art of record indicates that there are many problems to be overcome before all vector systems including retroviral vectors effect a contribution to medicine. Next, the artisan would look to the specification for guidance as to which compositions among the disclosed compositions, e.g., genetically modified cells expressing any protein, receptor, or enzyme for use in for the intended purpose of achieving a therapeutic effect as the result of protein drug expression, the specification provides little guidance for one skilled in the art to determine, without undue experimentation, as to which of the genetically modified stromal stem cell exhibit the intended "protein drug" effect in any animal having a disease, disorder, or condition as contemplated by the as-field specification.

More specifically with respect to claims directed enzyme therapy in the context of *ex vivo* gene therapy, even after 10 years from the effective filing date, Anderson (Nature, Vol. 292, 25-30, 1998) with respect to an experiment study on human ADA patients wherein autologous T cells are employed as carriers of ADA transduced retroviral vectors states that "although both girls have gene-engineered T lymphocytes in their circulation after more than 7 years, no definitive conclusion after more than 7 years, no definitive conclusion can be drawn as to the relative roles of PEG-ADA and gene therapy in their excellent clinical course". The fact that no working examples have been shown by the as-filed specification to conclusively show a therapeutically relevant effect in an animal having a real world medical condition associated with a protein defect, coupled with the unpredictability of gene therapy as expressed by the art of record, does not provide sufficient evidence for a skilled artisan to reasonably extrapolate, without any undue experimentation, from the basis of applicant's disclosure to any therapeutically useful effect by using any protein encoded vector contained in any bone marrow stromal cell encapsulated by any container as contemplated by the claimed invention.

While the as-filed specification indicates that by using a container known in the prior art, e.g., diffusion chambers, polymeric capsule, the genetically modified bone marrow stromal cells will be insulated or physically isolated from an immune response, thereby contemplating that a therapeutically relevant

Art Unit: 1633

effect can be generated by the cells due to the protection of the administered stromal stem cells by any container from the immune response of the treated animal. However, in addition to *in vivo* transient gene expression by an expression vector and the destruction of the diffused stromal stem cells by the immune response before the cells produce a sufficient amount of protein at a desired target site to produce a therapeutically relevant effect, Riddell *et al.* (Nature Medicine, Vol. 2, 2:216-223, 1996) reaffirmed the unpredictability of *ex vivo* gene therapy methods wherein foreign cells and/or proteins are employed at the time the invention was made by indicating that even with the use of autologous cells expressing a foreign protein, HIV-infected patients when grafted with autologous cytotoxic CD8+ T cells induce strong primary T-cell immune responses to foreign antigens expressed by transferred autologous cytotoxic CD8+ T cells (p. 221, column 1), and that the rejection of genetically modified cells and/or foreign proteins by even immunocompromised hosts suggests that *ex vivo* gene therapy by using foreign protein expressed transiently by even autologous cells is not routine or conventional in the prior art at the time the invention was made.

In addition, it is not clear how the systemic administration of genetically modified bone marrow cells which harbor a foreign receptor on the cell surface, for example, would not be destroyed by the immune response that is mounted against the foreign receptor expressed on the surface of the genetically modified cells after their diffusion from the container. Even if some of the genetically modified stromal stem cells and/or recombinant proteins escape from the immune response in a survived host after an administration or implantation, it is further not apparent how the genetically modified stromal stem cells traverse through barriers such as peripheral vein and endothelial wall to reach a disease site so as to generate a therapeutically relevant effect. Note also that Hoebe *et al.* (Human Gene Therapy, 4, 179-186, 1993) provides factual evidence showing that even if the genetically modified implanted fibroblast cells expressing a therapeutic protein, e.g., factor VIII, survive the immune response of a recipient mouse which is immunodeficient, there is no evidence of any recombinant Factor VIII in the plasma samples of recipient mice (abstract). The absence and/or *in vivo* transient expression of a recombinant protein in even a small animal such as immuno-deficient mice, and a rapid clearing of the introduced recombinant protein from mouse's



Art Unit: 1633

serum as shown in the prior art of record, does not lend any credible evidence to support applicant's claim that any stromal stem cell when carrying a protein drug encoded vector can be employed as a stable bioreactor to provide a continuous supply of protein drugs in any animal having any disease or disorder, such that a therapeutically relevant effect can be generated. Thus, where *ex vivo* gene therapy using any coding sequence of any protein is not reasonably predictable in establishing a therapeutic outcome of gene therapy for all types diseases, the gene therapy methods referred to in the present claims are also not predictable, nor is it apparent as to how a simple expression of endogenous protein from non-genetically modified MSC in transgenic mice as exemplified in the specification is reasonably correlated to a therapeutic effect in any animal having any disease as claimed.

Furthermore, the state of the art exemplified by Moritz *et al.* indicates that *ex vivo* gene therapy using genetically modified cell for engraftment into any an animals remain unpredictable. More specifically, the Moritz *et al.* reference (J. Clin. Invest. 1994, 93:1451-1457) indicating that "although gene transfer and long term gene expression in repopulating stem cells have been achieved in murine models by a number of investigators, *in vivo* experiments in larger animals such as dogs and primates have met with limited success, largely because of the low efficiency of infection of primitive hematopoietic stem cells"

The art of record clearly indicates that the efficiency of gene transfer into stem cells was much lower when similar protocols were assessed in large mammals and in human gene therapy trials, suggesting that there are species-specific differences in the susceptibility of stem cells to retrovirus infection. There are no working examples in the specification which indicates the efficiency of vector transduction in any stromal stem cell of any mammal including humans wherein a therapeutic effect is generated. Thus, in absence of any *in vivo* data regarding the grafting methods in any and/or all animals other than a murine model, it is not apparent how one skilled in the art determines the appropriate combination of transfection method, level of expression, cell numbers and method of administration for each possible gene, so as to have a therapeutic effect in any and/or all animals, without undue experimentation.

Art Unit: 1633

More specifically as to the state of the art of *ex vivo* gene therapy of employing any genetically modified hematopoietic cell expressing a transgene coding for an Ob protein, Onodera *et al.*, *Acta Haematologica*, 101, 2, pp. 89-96, 1999, indicates that even in 1999, the retroviral-mediated gene transfer to hematopoietic stems was insufficient for achievement of any therapeutically relevant effect (abstract). Kohn, *Current Opinion in Pediatrics*, 7, 56-63, 1995, indicates that the efficiency of gene transfer into stem cells was much lower when similar protocols were assessed in large mammals and in human gene therapy trials, suggesting that there are species-specific differences in the susceptibility of stem cells to retrovirus infection" (p. 58, column 1). Kohn further teaches that effective gene therapy for hematologic disorders remains unpredictable, and that a detection of circulating vector sequences in the blood *in vivo* after a transplantation of hematopoietic stem cells containing gene therapy vectors is not equivalent to a therapeutic effect (page 59, columns 1 and 2). There are no working examples in the specification which indicates the efficiency of transduction in bone marrow stromal or stromal stem cells of any mammal including humans wherein a therapeutic effect is generated.

Thus, it is clear from the evidence of record that that *ex vivo* gene therapy as claimed is not reasonably predictive and not yet shown to be successful. Applicants have not provided any convincing evidence that their claimed invention is indeed useful as a therapeutic for the treatment of any disease, disorder or condition as listed or contemplated by the as-filed specification, and have not provided sufficient guidance to allow one skilled in the practice the claimed invention without undue experimentation for the contemplated breadth of the claims. In absence of such guidance and evidence, the specification fails to provide an enabling disclosure.

In view of the lack of guidance regarding the administration parameters, lack of working examples, breadth of the claims, state of the art and the unpredictability of the art, as set forth by the evidence presented above, undue experimentation would be required by one of ordinary skill to practice the invention in the context of therapeutic application of *ex vivo* gene therapy as claimed.

The following is a quotation of the second paragraph of 35 U.S.C. 112:

Art Unit: 1633

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 72-75 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

The claims are indefinite in the recitation of "The container of claim 69, wherein the container is implanted in an animal" because it is not apparent as to how an active step of implanting the container of claim 69 in an animal is structurally contained in the container, nor is it apparent as whether or not applicant intends to claim an animal comprising the container of claim 69.

To the extent that claims 72-75 are interpreted as the container of claim 69 that is used for implantation in animal or a human, the following rejection is applicable.

***Claim Rejections - 35 U.S.C. § 103***

The following is a quotation of 35 U.S.C. § 103 which forms the basis for all obviousness rejections set forth in this Office action:

A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made. Subject matter developed by another person, which qualifies as prior art only under subsection (f) or (g) of section 102 of this title, shall not preclude patentability under this section where the subject matter and the claimed invention were, at the time the invention was made, owned by the same person or subject to an obligation of assignment to the same person.

Claims 69-75, 99-101, 106-108 and 112 are rejected under 35 USC 102(e) as being unpatentable over Carter *et al.* or Cerami *et al.*, each of which taken with any of Caplan *et al.* (US Pat No. 5,197,985), Schinstin *et al.* (US Pat No. 5,843,431), Mardon (previous cited prior art), and as evidenced by applicant's admission of the prior art of record as indicated on page 6 bridging page 7, and page 27 of the specification.

As reiterated from the previous Office actions, Carter *et al.* and Cerami *et al.* teach a method of

Art Unit: 1633

using isolated bone marrow stromal cells for implantation and providing a protein of interest to a cell in a mammal, wherein the cells are genetically modified cells to express a recombinant protein of interest (entire documents for Carter and Cerami *et al.* and also the teaching of implantation of isolated-genetically modified mesenchymal stem cells in Cerami *et al.*, column 6, lines 65-67, column 7, lines 1-67 and claim 1, column 8, lines 1-2). In addition, Cerami *et al.* also teach that the vector employed in the isolated mesenchymal stem cells can have regulatory and screening elements which include promoters and coding sequence of a selectable marker protein (column 7, lines 1-67).

Carter and Cerami *et al.* do not teach the use of a microcarrier, diffusion chamber, or microcapsule to control and/or enhance the delivery and release of the isolated bone marrow stromal cells.

However, at the time the invention was made, the prior art of record, as exemplified by Caplan, Schinstin *et al.*, and Mardon, does teach that it is routine and conventional to use a microcarrier, diffusion chamber, or microcapsule to enhance the delivery and subsequent release and differentiation of the implantable mesenchymal stem cells to the target site (see entire document of each of the cited reference).

Furthermore, the specification teaches on page 3 bridging page 7 that immunological isolation means include well known technologies and devices such as microencapsulation, diffusion chambers, etc.

It would have been obvious for one of ordinary skill in the art to have employed any known a microcarrier, diffusion chamber, or microcapsule to enhance the delivery and subsequent release of the implantable mesenchymal stem cells disclosed in Carter or Cerami *et al.* to the target site. One of ordinary skill in the art would have been motivated to have employed a microcarrier, diffusion chamber, or microcapsule to enhance the delivery and subsequent release of the implantable mesenchymal stem cells to the target site because of the reasons set forth in the immediately preceding paragraph. Note that use of well known technologies and devices such as microencapsulation, diffusion chambers, etc., as taught by the combined cited references, would physically isolate the isolated genetically modified stromal cells from the immune response, as evidenced by applicant's admission of the prior art of record as indicated on page 6 bridging page 7 of the specification.

In addition, one of ordinary skill in the art would have been motivated to transfect or transduce the

Art Unit: 1633

cells of Carter by conventional methods with vectors containing any known promoter, signal sequences, beneficial protein, and/or a coding sequence of a selectable marker, such as those disclosed in the cited references to determine and track the effect of these regulatory elements and the subsequent expression of a desired gene during the differentiation of the stromal cells once implanted in an animal model, as disclosed in Carter and Cerami *et al.* references.

It would also have been obvious for one of ordinary skill in the art to have employed any pore size in any of the containers available in the prior art of record as an obvious matter of design choice particularly since such modifications would be expected to lead to an equivalent enhancement in delivery, release, expression and differentiation of the cells at the target delivery site, particularly in view of the absence of factual evidence showing an unexpected property of the use of the claimed pore size relative to those outside the claimed diameter of the pores of the claimed containers.

Therefore, the invention as a whole is prima facie obvious, as evidenced by the references, especially in the absence of evidence to the contrary.

Claims 69 and 77 are rejected under 35 USC 102(e) as being unpatentable over Carter *et al.* or Cerami *et al.*, each of which taken with any of Caplan *et al.* (US Pat No. 5,197,985), Schinstin *et al.* (US Pat No. 5,843,431), Mardon (previous cited prior art), and as evidenced by applicant's admission of the prior art of record as indicated on page 6 bridging page 7 of the specification, and further in view of Beresford *et al.* and Flier (cited in the previous office actions).

The rejection of the base claim 29 as being unpatentable over Carter *et al.* or Cerami, each of which taken with any of Caplan *et al.*, Schinstin *et al.*, Mardon (previous cited prior art), and as evidenced by applicant's admission of the prior art of record, is applied here as indicated above.

The combined cited references do not teach that the recombinant protein encoded DNA is the obesity factor encoded DNA.

However, at the time the invention was made, Beresford *et al.* disclose that rat marrow stromal cell cultures are capable of differentiating into adipocytic and osteogenic cells, and further, are capable of

Art Unit: 1633

expressing collagens (type I, III, and IV). The relative amounts of cells which differentiate into adipocytes and osteogenic cells, as well as the relative types and amounts of collagens synthesized by the cells is dependent of the culture conditions of the stromal cells (see page 344-345, under "Collagen synthesis", and page 348, left column, second paragraph). As the obesity factor encoded DNA is expressed in adipocytes (see Flier), one of ordinary skill in the art would have been motivated to generate a vector construct containing the obesity factor encoded DNA, operatively linked to regulatory elements associated with collagens types I, III and IV for the purpose of determining the effect of these regulatory elements on the expression of the obesity gene, and the effect of the obesity gene in stromal cells during differentiation into adipocytes or osteogenic cells. Moreover, as the references disclose transfecting stromal cells with expression vector constructs comprising a gene of interest, and in view of the teachings of Flier that the obesity protein is expressed in cells of the mesenchymal cells (stromal cells) lineage, one of ordinary skill in the art would have had a high expectation of successfully transfecting stromal cells with an expression vector encoding an obesity factor, such that the transfected cells synthesize the obesity factor, which has previously been established to occur in differentiated cells of the stromal cell lineage, barring evidence to the contrary.

Carter *et al.* or Cerami *et al.*, each of which taken with any of Caplan *et al.* (US Pat No. 5,197,985), Schinstin *et al.* (US Pat No. 5,843,431), Mardon (previous cited prior art), and as evidenced by applicant's admission of the prior art of record.

Therefore, the invention as a whole is prima facie obvious, as evidenced by the references, especially in the absence of evidence to the contrary.

Applicant's response (pages 8 and 9) has been considered by the examiner but is moot in view of the new grounds of rejection as set forth above.

No claims are allowed.

Art Unit: 1633

Any inquiry concerning this communication or earlier communications regarding the formalities should be directed to Patent Analyst Kimberly Davis, whose telephone number is **(703) 305-3015**.

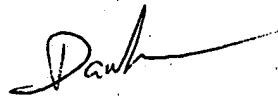
Any inquiry concerning this communication or earlier communications from the examiner should be directed to examiner *Dave Nguyen* whose telephone number is **(703) 305-2024**.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, *Deborah Clark*, may be reached at **(703) 305-4051**.

Papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Group 1600 via the PTO Fax Center located in Crystal Mall 1. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). The CM1 Fax Center number is **(703) 305-7401**.

Any inquiry of a general nature or relating to the status of this application should be directed to the *Group receptionist* whose telephone number is **(703) 308-0196**.

Dave Nguyen  
Primary Examiner  
Art Unit: 1633



**DAVE T. NGUYEN**  
**PRIMARY EXAMINER**